AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) A method of treating viral infections in a patient which method comprises co-administering to said patient a therapeutically effective amount of interferon and a low-dose of ribavirin, wherein at least said ribavirin is administered in a slow-release formulation at a dosage of less than 400 mg ribavirin per day to provide a clinically effective blood level in a portal vein and less than required to provide a clinically effective blood level in a peripheral circulation to thereby provide a dose-delivery rate having a selective antiviral and interferon potentiating effect in a liver of the patient.

Claims 2-4 (Canceled).

- 5. <u>(Currently Amended)</u> The method according to claim 41, wherein the controlledslow-release formulation releases ribavirin by a mechanism chosen from diffusion and erosion.
- (Currently Amended) The method according to claim 41, wherein the eentrolledslow-release formulation of ribavirin comprises at least one of polymer-coated multiparticulates, polymer-coated tablets, polymer-coated minitablets, and hydrophilic matrix tablets.

Claims 7-8 (Canceled).

- 9. (Currently Amended) A method according to claim 81, wherein the ribavirin dose is in the range of from 20 to 350 mg/day.
- (Original) A method according to claim 1, wherein the ribavirin dose is varied according to the body weight of the patient.
- (Original) A method according to claim 10, wherein the ribavirin dose is less than 6 mg/kg/day.
- (Original) A method according to claim 11, wherein the ribavirin dose is less than 5 mg/kg/day.

- (Original) A method according to claim 12, wherein the ribavirin dose is in the range of from 1 to 5 mg/kg/day.
- (Original) The method according to claim 13, wherein the viral infection is hepatitis C.
- 15. (Original) The method according to claim 1, wherein the ribavirin is in the form of at least one of a ribavirin ester, salt, or analogue of ribavirin shown to be effective as an antiviral agent.
- (Original) The method according to claim 15, wherein the interferon is interferon alfa or pegylated interferon alfa.
 - 17. (Original) The method of claim 16, wherein the interferon is interferon alfa 2b.
- 18. (Original) The method according to claim 17, wherein the interferon is administered parenterally.
- 19. (Original) The method according to claim 18, wherein the interferon is administered by subcutaneous IV or IM injection.
- 20. (Original) The method according to claim 19, wherein the interferon is administered parenterally in an amount of from 2 to 10 million IU per week on a weekly, thrice weekly ("TIW"), every other day ("OOD") or daily basis.
- 21. (Original) The method according to claim 16, wherein the pegylated interferon alfa is pegylated interferon alfa-2b and is administered systemically in an amount of 0.5 to 2.0 micrograms per kilogram of body weight per week on a weekly, TIW, QOD or daily basis.
- 22. (Original) A method according to claim 16, wherein the pegylated interferon alfa is pegylated interferon alfa-2a and is administered systemically in an amount of 20 to 250 micrograms per kilogram of body weight per week on a weekly, TIW, QOD or daily basis.
- 23. (Currently Amended) A method of treating viral infections in a patient which comprises co-administering to said patient a therapeutically effective amount of interferon with less than 400 mg/day of ribavirin which is administered as a slow release formulation.

Claims 24-26 (Canceled).

 (Currently Amended) A method according to claim <u>2326</u>, wherein the ribavirin dese-is <u>provided</u> in <u>a dosage</u>the range of from 5 to 399 mg/day. Application No. 10/568,176 Amendment Dated 7/21/2008 Reply to Office Action of 3/20/2008

- 28. (Canceled).
- (Currently Amended) A method according to claim <u>2328</u> further comprising <u>administering systemic doses of an antioxidant or other membrane protective agent—which is administered in systemic doses.</u>
- (Currently Amended) A method according to claim <u>2328</u> further comprising administering an antioxidant or other membrane protective agent which is administered as a lowdose, slow-release formulation.
- (Currently Amended) A method according to claim <u>2328</u>, further comprising <u>administering</u> an antioxidant or other membrane protective agent which is co-formulated with the ribavirin as a low-dose-slow-release formulation.
 - 32. (Canceled)
- 33. (Currently Amended) A kit for use in the treatment of viral infections comprising a therapeutically effective amount of interferon in combination with ribavirin and optionally an antioxidant or other membrane protective agent as a slow-release formulation comprising less than 400 mg per day of ribavirin.

Claims 34-35 (Canceled).

- . 36. (Currently Amended) A kit according to claim 3335 wherein the slow-release formulation of ribavirin comprises at least one of polymer-coated multiparticulates, polymer-coated tablets, polymer-coated minitablets, and hydrophilic matrix tablets.
 - 37. (Canceled).
- 38. (Currently Amended) A kit according to claim 3335 wherein athe unit dose of ribavirin is less than 6 mg/kg/day.
- 39. (Original) A kit according to claim 33 wherein the ribavirin is in the form of at least one of a ribavirin ester, salt or analogue or ribavirin shown to be effective as an antiviral agent.
- 40. (Original) A kit according to claim 33 wherein the interferon is in a form for parenteral administration.
- 41. (Original) A kit according to claim 33 comprising unit doses of interferon for providing an amount of from 2 to 10 million IU per week by thrice weekly ("TIW"), every other day ("QOD") or daily administration.

Application No. 10/568,176 Amendment Dated 7/21/2008 Reply to Office Action of 3/20/2008

- 42. (Original) A kit according to claim 33 wherein the interferon is interferon alfa or pegylated interferon alfa.
- 43. (Currently Amended) A pharmaceutical composition for the treatment of viral infections in a patient comprising a therapeutically effective amount of interferon together with a lew-dose of ribavirin and optionally an antioxidant or other membrane protective agent, wherein a daily dosage amount of the pharmaceutical composition contains less than 400 mg of ribavirin.